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cells are provided in an amount effective to inhibit smooth muscle cell proliferation at the

Vagentar endonettal cell of the licit vessel

site of the injury without migration of theiendothelial cells to the enterior lining.

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3. (twice amended) The method of claim 1 wherein the matrix is in a form selected from the group consisting of gels [or], foams, suspensions, microcapsules, solid polymeric supports, [or] and fibrous structures.

A)

6. (twice amended) The method of claim 5 wherein the matrix is formed of a material selected from the group consisting of polyhydroxy acids, polyorthoesters, polyanhydrides, proteins, carbohydrates [or] \_\_and\_ polysaccharides, polyphosphazenes, polyalkylene oxides and combinations thereof.

8. (twice amended) The method of claim 1 wherein the matrix further comprises biologically active compounds selected from the group consisting of [,] prostaglandins, prostanoids, [compounds regulating the renin-angiotensin axis,] tyrosine kinase inhibitors, immunosuppressants, glucocorticoids, anti-oxidants, free radical scavengers, peptide hormones, angiogenic and angiogenic inhibitory factors, and combinations thereof.

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11. (twice amended) A composition for inhibiting mechanisms involved in having an enouteful cell lining the enouteful cell lining restensis of a blood vessel/following injury to vascular tissue/of the blood vessel in a patient in need of treatment thereof, comprising a biocompatible matrix shaped for implantation adjacent to a blood vessel, the matrix having seeded therein or thereon dissociated/endothelial vascular cells, wherein the/endothelial cells are in an amount effective to inhibit smooth muscle cell

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proliferation at the site of the injury without migration of theyendothelial cells to the material.

of the blood vessel

13. (twice amended) The composition of claim 11 wherein the matrix is in a form selected from the group consisting of gels [or], foams, suspensions, microcapsules, solid polymeric supports, [or] and fibrous structures.

14. (amended) The composition of claim 11 wherein the cells are selected from the group consisting of autologous cells, allograft cells, and xenograft cells[, and genetically engineered cells].

16. (twice amended) The composition of claim 15 wherein the matrix is formed of a material selected from the group consisting of polyhydroxy acids, polyorthoesters, polyanhydrides, proteins, earbohydrates [or], polysaccharides, polyphosphazenes, and combinations thereof.

18. (twice amended) The composition of claim 11 wherein the matrix further comprises biologically active compounds selected from the group consisting of prostaglandins, prostanoids, [compounds regulating the renin-angiotensin axis,] tyrosine kinase inhibitors, immunosuppressants, glucocorticoids, anti-oxidants, free radical scavengers, peptide hormones, angiogenic and angiogenic inhibitory factors.

Please add new claim 20.

(new) The composition of claim 14 wherein the cells are genetically engineered.

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